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Modulation of gene expression via overlapping binding sites exerted by ZNF143, Notch1 and THAP11

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ABSTRACT

ZNF143 is a zinc-finger protein involved in the transcriptional regulation of both coding and non-coding genes from polymerase II and III promoters. Our study deciphers the genome-wide regulatory role of ZNF143 in relation with the two previously unrelated transcription factors Notch1/ICN1 and thanatos-associated protein 11 (THAP11) in several human and murine cells. We show that two distinct motifs, SBS1 and SBS2, are associated to ZNF143-binding events in promoters of >3000 genes. Without co-occupation, these sites are also bound by Notch1/ICN1 in T-lymphoblastic leukaemia cells as well as by THAP11, a factor involved in self-renewal of embryonic stem cells. We present evidence that ICN1 binding overlaps with ZNF143 binding events at the SBS1 and SBS2 motifs, whereas the overlap occurs only at SBS2 for THAP11. We demonstrate that the three factors modulate expression of common target genes through the mutually exclusive occupation of overlapping binding sites. The model we propose predicts that the binding competition between the three factors controls biological processes such as rapid cell growth of both neoplastic and stem cells. Overall, our study establishes a novel relationship between ZNF143, THAP11 and ICN1 and reveals important insights into ZNF143-mediated gene regulation.

INTRODUCTION

The transcriptional regulatory system plays a fundamental role in controlling the correct expression of genes involved in many biological processes (1). This mechanism involves specific DNA-binding transcription factors, co-factors and chromatin remodelling factors. In humans, there are nearly 1400 transcription factors of which only a few have been extensively studied (1). They bind to cis-regulatory elements located in the promoters of specific genes and modulate their activation or repression (2). Therefore, their influence on gene expression is achieved by acting directly, or via a multiplicity of partners, on chromatin remodelling and recruitment of the transcription machinery (3). Despite our limited knowledge of the function of all transcription factors, cis-regulatory regions of genes are broadly studied nowadays at the genome-wide scale (4). Owing to the increasing number of high-throughput studies (5,6), numerous transcription factors are found to be located within the same promoter regions affecting or not the gene expression, acting in synergy or having antagonistic effects depending on physiological and environmental conditions (7,8). Thereby combinatorial binding of transcription factors modulates gene expression according to the needs and conditions of cells (9).

In this regard, we were first interested in deciphering the genome-wide regulatory potential of the transcription factor ZNF143. Also known as Staf (Seleno cysteine tRNA gene transcription activating factor), it is a zinc-finger protein involved in the control of both coding and non-coding genes from RNA polymerase II (Pol II) and RNA polymerase III (Pol III) promoters (10). This factor recognizes and binds with high affinity a well-characterized 18-bp motif located in the core promoter region. ZNF143 has been shown to be involved in the transcriptional regulation of DNA repair genes (10–12) and protein-coding genes (13–15). It is conserved in all chordates, and its vertebrate parologue ZNF76 (16) possesses an identical central 209 amino acid long DNA-binding domain (DBD). ZNF143 is involved in the resistance to cis-platin in cancer cells through the transcriptional regulation of DNA repair genes (17). Moreover, this factor is critical for the normal development of zebrafish embryo...
Antibodies and constructs used in these studies are described in Supplementary Materials and Methods (Supplementary data).

**MATERIALS AND METHODS**

**Antibodies and constructs**

Antibodies and constructs used in these studies are described in Supplementary Materials and Methods (Supplementary data).

**Generation of stable cell lines and induced protein expression**

F荧光素TM T-Rex™ 293系统（Invitrogen）用于根据制造者的说明来生成稳定细胞系，并且使用调节表达ZNF143 (FLP143)，ZNF143-3xHA（FLP143-HA），ZNF76 (FLP76) RBPJ-3xHA（FLP_RBPJ-HA）和THAP11-3xHA（FLPTHAP11-HA）的稳定克隆。这些克隆携带了所需的开放阅读框架，这些开放阅读框架在控制细胞中表达的转录因子中得到了重复的读取。已经使用诱导的蛋白质表达来检测doxycycline regulation protein expression。蛋白质表达是通过添加1 μg/ml的doxycycline来获得的。

**ChiP and ChiP-Seq**

ChiP assays were performed essentially as previously described (28) with 1–4 × 10⁷ cells. Chromatin was sonicated to an average of 200 bp using a Bioruptor™ UCD-200 (Diagenode) at maximum power level for 30 min (pulses of 30 s with 30 s breaks). After DNA purification, enrichment was estimated by quantitative real time PCR (qPCR). qPCR was performed in duplicate on a Stratagene MX3005P PCR system (Agilent Technologies) using 5× HOUT Pol® EvaGreen® qPCR Mix PLUS (ROX) (Euromedex) and input DNA as the standard. Promoters containing, and regions devoid of the studied binding sites, were used as positive and negative controls, respectively. Enrichment was determined by the ΔΔCt method. Primer sequences are available on request. Samples were sequenced using the Illumina Genome Analyzer GAII, and the read sequences were mapped to unmasked version of the human (hg19) or mouse (mm9) genomes via the Illumina Eland V1.6 (Illumina) or by the Bowtie software (29) with either an exact 36 nt match or with a single or two mismatches. Reads mapped to multiple locations were discarded, and no more than two redundant reads having exactly the same genomic coordinates were used to identify enriched regions. Tag mapping in the large families of highly related ncRNA genes was performed with the Bowtie software, allowing two perfect matches in the genome. ChiP-Seq data were submitted to the Gene Expression Omnibus database and assigned the identifier GSE39263.

Description of ChiP-Seq data processing and integrative analysis is available in Supplementary Materials and Methods (Supplementary data).

**Electrophoretic mobility shift assay**

Recombinant ZNF143 DBD, RBPJ from VK91 (30) and THAP11 DBD were produced using the GST gene fusion system as previously described (15,30) essentially according to the manufacturer’s instructions (GE Healthcare). Dissociation constants of ZNF143 on the SBS1 (GTTA TGGATTTCCATTATGCACCGC) or SBS2 (AAAC TACAATTCCCCATATGCGC) motifs were derived as previously described (16). Internally labelled

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DNA fragments containing the SBS1 (TTCCCATTATG CACCGG), SBS2 (ACTACAATTCCATTAGCAC CGCG), THAP11-binding site (TBS) (ACTACAATT CACCGG), and ACTACAATT CCCA) and ACTACAAT complexes were resolved by electrophoresis on 4% native polyacrylamide gels containing 0.25 × Tris-borate-EDTA. Competition was performed with a 200-fold molar excess of unlabelled specific competitor or a unspecific competitor.

RNA preparation and expression analysis

ZNF143 knockdown was performed in HeLa cells as described previously (28). The THAP11-specific Mission esiRNA (Sigma-Aldrich) targeting positions 872–1276 of the THAP11 mRNA or control, esiRNA targeting GFP (Sigma-Aldrich) were transfected into HeLa cells using Lipofectamine2000 according to manufacturer’s instructions (Invitrogen). For γ-secretase inhibition, HB-P-ALL cells were treated with 500 nM of Compound E (Enzo Life Sciences), and control was obtained with DMSO-treated HB-P-ALL cells. ZNF143-HA or ZNF143 overexpression was induced by treatment of FLP143-HA or FLP143 cells with 1 µg/ml of doxycycline, and cells were collected at different time points post-induction. Total RNA was extracted using TRIAGENT (Euromedex) and reverse transcribed with dN9 oligonucleotides. The complementary DNA obtained was amplified with specific primers on the Stratagene Mx3005P PCR system (Agilent Technologies) using EvaGreen qPCR Mix Plus (Euromedex). Primer sequences are available on request. All reactions were carried out in triplicate. The relative expression ratio was calculated using the REST 2009 software (Qiagen).

RNA-seq differential expression analysis

Total RNA was isolated from HeLa-S3 and FLP143 cells treated as described earlier in the text. After selection of polyA+mRNA, RNA-seq libraries preparation was performed using Illumina TruSeq RNA Sample Preparation kit according to manufacturer protocol. All RNA samples were sequenced using the Illumina Genome Analyzer GAII and aligned to the hg19 reference genome using TopHat software (31). Aligned reads were processed with Cuffdiff (32) to measure the differential expression using the RefSeq genes as reference annotation. Only the transcripts matching the genes potentially regulated by ZNF143 (TSS located at +/−2 kb of a ChIP-Seq peak summit) were analysed, and the expression value for each transcript was measured by a FPKM value ( Fragments per kilobase of transcripts per million mapped tags). RNA-seq data were submitted to the Gene Expression Omnibus database and assigned the identifier GSE39263.

Data access

Data from this study were submitted to the NCBI Gene Expression Omnibus (http://www.ncbi.nlm.nih.gov/geo) under accession number GSE39263.

RESULTS

Genome-wide identification, characterization and distribution of ZNF143-binding events in different human and mouse cell lines

To identify ZNF143-binding events, ChIP-Seq was performed with anti-ZNF143 antibody in four human (HeLa-S3, T-Rex HEK293, K562 and HPB-ALL; Supplementary Figure S1A) and three mouse cell types (NIH3T3, mESC and MEF; Supplementary Figure S1B). Quantitative enrichment analysis of the ZNF143 peaks showed a strong correlation (coefficient between 0.77 and 0.93) between the experiments (Figure 1A and Supplementary Figure S2). The locations of ZNF143-binding events show an important enrichment in peaks located at +/−2 kb from the transcription start site (TSS) of protein coding genes (50 and 59% of all human and mouse ZNF143-binding events, respectively) (Supplementary Table S1) (Figure 1B and C and Supplementary Figure S3A). In human and mouse cells, we identified the MLT11 retrotransposon subfamily, showing a significant enrichment in ZNF143 peaks compared with random coordinates (88.9-fold enrichment in HeLa cells). MLT11 retroelements associated with ZNF143 are essentially located outside of promoters and represent 35.5% of repeat elements overlapping ZNF143-binding events in HeLa cells (Figure 1D). We will focus now on ZNF143-binding events located and enriched at proximity of the TSS of coding genes (+/−2 kb from a TSS) (Supplementary Figure S3B). We identified, at these locations, 3122 ZNF143-binding events in the set of human (Supplementary Table S4) and 3385 in the set of mouse cells (Supplementary Table S5), 60.3% of which being conserved in humans (Figure 1E). We observed a core set of 2504 ZNF143-binding regions shared by the four human cell lines (Figure 1F) and one of 2686 shared by the three mouse cell types (data not shown). The binding events were essentially present in the 500 bp upstream of the TSS (Supplementary Figure S4) with a positional preference at 70 bp upstream of it (Figure 1G). We observed also an important enrichment of the ZNF143-binding events located in bidirectional promoters. In human cells, 661 binding events are located within bidirectional promoters (Supplementary Table S4), representing 17% of all the bidirectional promoters in the genome. The chromosomal distribution of the ZNF143-binding events showed a noteworthy enrichment of the ZNF143 peaks in promoters of genes encoding krüppel-associated box (KRAB) zinc-finger proteins on the human chromosome 19 and mouse chromosome 7 (Supplementary Figure S5). Finally, we examined the ZNF143 occupancy in five sets of ncRNA genes (RefSeq ncrNA, ensembl ncrNA, lincRNA, TUCP and Pol III ncRNA). We identified 476
Figure 1. Genome-wide characterization of ZNF143-binding events in human and mouse cells. (A) Correlation plots of ZNF143-binding events in four human cell lines. Grid of pairwise scatter plots with associated Pearson correlation coefficient obtained using Cistrome software (33). The blue dots coordinates represent the normalized enrichments in reads for each cell lines at all the peaks coordinates (Supplementary Table S1). The regression line is indicated in red and the Pearson correlation coefficients are indicated in the squares on the top right. (B) Pie charts showing distribution of the ZNF143-binding events in different genomic regions. A promoter is defined as a region within 2 kb from the TSS of a gene; downstream, 2 kb downstream of the end of a gene; the distal intergenic are regions excluding a promoter, downstream, 5'UTR, 3'UTR, introns and

(continued)
Two distinct motifs with overlapping sequences are associated to the in vivo ZNF143-binding events

We identified by a motif search conducted on all ZNF143-binding events the known 18-bp ZNF143-binding motif (10) and a second 25-bp motif containing the first one with an additional ACTACAN sequence located immediately upstream (Figure 2A). Henceforth, we will refer to the canonical 18-bp motif as SBS1 and the 25-bp motif as SBS2 (Figure 2A). The ZNF143-binding events that overlap MLT1J retroelements contain only the SBS2 motif (Figure 2B). For motifs associated to the ZNF143 peaks identified at proximity of ncRNA genes, we determined that the binding events in subsets 1 and 2 (Supplementary Figure S6) include the SBS1 and/or SBS2 motifs. Surprisingly, all the 56 ZNF143-binding events associated to Pol III type 3 ncRNA and Pol II snRNA genes (subsets 3 and 4 in Supplementary Figure S6) contain only one SBS1 motif while we would expect only 10 in respect of the distribution of SBS1 and SBS2 motifs in the all binding events (z-test: \( P < 10^{-5} \)). Subset 3 contains peaks located at proximity of the TRNAU1 (tRNAsec), RNU6.1, RNU6.2, RNAU6.7-U6.9, RN7SK, RPPH1, RNY1, RNY3, RNY4 and RNY5 genes (Supplementary Table S7). Supplementary Figure S7A (lower part) displays a multiple alignment showing sequences of the four elements (TATA element, PSE promoter element, octamer motif and SBS1) (35) constituting the promoters of the 12 genes cited earlier in the text. The position of the SBS1 motif is restricted to a window covering positions \(-245\) to \(-199\), except for the promoter of the RPPH1 gene (11) where SBS1 is located at position \(-87\). Subset 4 (Pol II snRNA genes) contains the RNU1, RNU2, RNU4, RNU4ATAC, RNU5, RNU6, RNU11 and RNU12 genes (Supplementary Table S7). The multiple sequence alignment in Supplementary Figures S7A (upper part) and S7B shows the presence of canonical PSE and octamer motifs at proximity of SBS1 in 41 of the 44 Pol II snRNA genes (Supplementary Table S7). We will henceforth focus on the SBS1 and SBS2 motifs identified in promoters (+/-2 kb from a TSS) of protein coding genes in humans. The SBS1 and/or SBS2 motifs were detected in 74% of the 3122 binding events (Figure 2C and Supplementary Table S4). As ZNF143 could be targeted to certain locations by some of its putative partners, we searched for known binding sites in the peaks without any obvious ZNF143-binding motif. The only other motif reported was an Sp1-like binding site; it was enriched around the peaks, reflecting per se the association of ZNF143 with CpG islands. Figure 2C shows that the family of peaks with SBS1 and/or SBS2 motifs consists of three subfamilies: one subfamily (32% of all peaks) contains only SBS1 motifs, the second (23%) only SBS2 motifs and the third one (19%) both motifs. We observed that 21 and 14% of all binding events contain only a single SBS1 or SBS2 motif, respectively (Figure 2C). Strikingly, in contrast to the binding events with only an SBS1 or an increasing number of SBS1 motif, the ZNF143-binding events with a single or increasing number of SBS2 motifs showed a increasing ChIP score (t-test: \( P < 10^{-5} \)) (Figure 2D and Supplementary Figure S8) Finally, we examined the functional annotation of genes at proximity of the 2504 core set of ZNF143-binding events shared by the four human cell lines (Figure 3A). We compared then four sets of genes identified at proximity of the peaks and containing different combinations of motifs (SBS1 motif only, SBS2 motif only, both motifs) or without motif (Supplementary Figure S9). Overall, the functional analysis revealed that all the sets of genes are enriched in terms associated to cell growth and primary metabolism. Nevertheless, a high enrichment score was found for terms such as translation for the set of genes associated to SBS1 and transcription regulation and zinc fingers for the SBS2 motif. To attest the involvement of ZNF143 in cell growth and primary metabolism, we tested the impact of ZNF143 knockdown on HeLa-S3 cells proliferation. As expected and as previously shown for PC3 cells (37), HeLa-S3 cells had a significantly reduced proliferation rate at 72 h post-transfection with siRNA targeting specifically ZNF143 versus siRNA-targeting GFP (Figure 3B).
transfected with siRNA-targeting ZNF143 or GFP. Results are expressed as means ± standard deviation of three biological replicates.

**Figure 2.** Identification and characterization of ZNF143-binding motifs. (A) Sequence logo depicting the SBS1 and SBS2 motifs discovered de novo at +/−100 bp ZNF143 peak summits in human and mouse genomes, using the MEME suite (34). (B) Multiple sequence alignment depicting some of the MLT1J elements that are bound by ZNF143. Coordinates of the repeated elements are shown on the left with indication of the orientation of the element (+/−). (C) ZNF143-binding event counts in human genome associated with SBS1, SBS2, SBS1 and SBS2, a single SBS1 or a single SBS2 motif. (D) Box plots showing ChIP-Seq score (number of reads) distribution of ZNF143-binding events containing: only SBS1 or SBS2 motifs.

Genome-wide analysis of ZNF76 binding in T-REx HEK293 cells reveals a total overlap with ZNF143 binding

In mammalian cells, ZNF76 was characterized as the ZNF143 parologue with an activator and repressor function (16,38). We asked whether the identified ZNF143 targets are recognized by ZNF76 in human T-Rex HEK293 cells expressing ZNF76 (Supplementary Figure S1D). To facilitate ChIP, clonal cell lines expressing ZNF76 were generated (Supplementary Figure S1A), and we characterized the loci occupied by this factor by ChIP-Seq. The data were compared with those obtained by the anti-HA ChIP-Seq on FLP143-HA cells. An enrichment correlation coefficient of 0.94 was observed between the two experiments for all of the binding events (Figure 1H). Obviously, a de novo motif search showed that the ZNF76-binding events include the SBS1 or SBS2 motifs described earlier in the text. Moreover, both ZNF76 and ZNF143 behave, in terms of enrichment on SBS1 or SBS2 sites, in an identical way. This is illustrated in Supplementary Figure S10 with the peaks formed on the ATP5J2 (SBS1 only) and NUP153

**Figure 3.** ZNF143-targeted genes are involved in cell growth and proliferation. (A) Functional categories enriched among the genes located at +/−2 kb of the identified ZNF143-binding events common to HeLa-S3, K562, T-Rex-293 and HPB-ALL cell line (Supplementary Table S4), as reported by the DAVID web-based functional annotation program (36). Values are fold-enrichment scores compared with the whole set of human genes used as background. (B) Cell proliferation assay performed for a 72 h time course on HeLa-S3 cells non-transfected (no siRNA) or transfected with siRNA-targeting ZNF143 or GFP. Results are expressed as means +/− standard deviation of three biological replicates.
(SBS2 only) promoters. We evidenced also that ZNF76 is associated to ncRNA genes with a total overlap with the ZNF143-binding events.

**THAP11 and ZNF143 bind genome-wide to SBS2 sites through mutually exclusive events**

The widespread association of SBS1 and/or SBS2 motifs to ZNF143-binding events led us to examine whether the additional ACTACAN sequence in SBS2 affects ZNF143 binding. The apparent Kd of ZNF143 for the SBS1 (1.36 ±/−0.33 nM) and SBS2 (1.58 ±/−0.19 nM) motifs are very similar (Supplementary Figure S11), suggesting that the 5' part of the SBS2 motif (ACTACAN), or a more extended sub-motif, constitutes a binding site for another factor. In this respect, ChIP-Seq on mES cells revealed that the transcription factor Thap11 binds with its transcriptional co-regulator Hcf-1 to the sequence motif ACTACANNCTCCA (TBS) (Figure 4A) (25), suggesting a possible overlap of ZNF143 and Thap11 DNA-binding events. We compared the available ChIP-Seq data of Thap11 and its co-factor Hcf-1 (25) obtained from mES cells with our ZNF143 ChIP-Seq data obtained from the same cells. We found that 84.7% of all Thap11 and 75.8% of all Hcf-1-binding events located in promoters overlap the ZNF143-binding events located at +/-2 kb of a TSS (3385 peaks) (Figure 4B). An individual motif occurrence search on the 651 peaks, common to the Thap11- and ZNF143-binding events (Figure 4B), identified an SBS2 motif in the whole set of coordinates. Furthermore, a similar search performed on Hcf-1 peaks without reported Thap11 binding, showed that 51% (Figure 4B) of them overlap with ZNF143 and contain also an SBS2 motif. This suggests that Thap11 could in fact also bind these SBS2 sites. From these observations, it appears that Thap11 binding is restricted to loci containing an SBS2 motif. By ChIP, we asked whether the promoters containing only SBS1 (SBS1 loci) or SBS2 (SBS2 loci) motifs are recognized by THAP11. We generated clonal cell lines expressing HA-Tagged THAP11 (FLPTHAP11-HA) and by anti-HA ChIP, we examined and compared the THAP11 and ZNF143 occupancy on SBS1 and SBS2 loci (Figure 4C). As expected, ChIP-Seq showed ZNF143 occupancy on the nine SBS1 loci and on the eight SBS2 loci (Figure 4C, lower part). In contrast, THAP11 occupancy was restricted to the SBS2 loci without significant occupation of the SBS1 loci (Figure 4C, upper part). By gel-shift experiments, we further examined the in vitro binding of ZNF143 and THAP11 DBDs on TBS, SBS1 and SBS2 motifs. As expected, the ZNF143-DBD binds specifically to the SBS1 and SBS2 motifs (Figure 4D, lanes 1–8) but is unable to recognize the TBS (Figure 4D, lanes 9 and 10). THAP11-DBD recognized specifically the TBS (Figure 4D, lanes 11–14) and the SBS2 motif (Figure 4D, lanes 15–18), which per se contains a TBS. Surprisingly, unlike the TBS sub-motif ACTACAA (Figure 4D, lanes 23–26), the SBS1 motif, which contains only the 3' part of TBS (TTCCCA) binds specifically the THAP11–DBD (Figure 4D, lanes 19–22). This observation suggests that THAP11 has a higher affinity to

the 3'-part of the TBS (TTCCCA). Binding assays performed on the SBS2 motif in the presence of ZNF143-DBD and THAP11–DBD revealed that the binding of the two proteins is mutually exclusive (Figure 4D, lanes 27–30). Indeed, the shift observed in the presence of THAP11 (Figure 4D, lane 29) is not altered by the presence of ZNF143–DBD (Figure 4D, lane 30). We also tested in vivo the THAP11 occupancy by ChIP on loci containing SBS2 motifs after ZNF143 siRNA-mediated knockdown in non-induced or induced FLPTHAP11-HA cells. The induced expression of THAP11 increases the occupancy of the factor on the promoters (Figure 4E, left part, compare induced and non-induced after siCTRL treatment). This phenomenon is amplified after THAP11 induction performed on cells with efficient ZNF143 siRNA-mediated knockdown (Figure 4E, left part, compare induced and non-induced after siRNA ZNF143). The efficiency of ZNF143 depletion was validated by anti-ZNF143 ChIP qPCR, as illustrated in Figure 4E (right part) for the AHSAT1 promoter.

Finally, we determined the over-represented functional categories of genes located near loci occupied by both ZNF143 and Thap11 (36). As for the whole set of promoters recognized by ZNF143, the enriched terms concern cell growth and primary metabolism, whereas cell signalling and cell development were under-represented. The association of ZNF143 to the MLT1J retroelement via a SBS2 motif suggests a possible association of THAP11 to the same retroelement. Indeed, we verified by anti-HA ChIP-qPCR that THAP11 is also associated in vivo to an MLT1J retroelement locus (Supplementary Figure S12A).

**Genome-wide analysis of Notch1/ICN1 binding in HPB-ALL cells reveals widespread overlap with ZNF143**

The ICN1 protein, the active form of the Notch1 receptor, is recruited to promoters by RBPJ, which binds to the TTCCCA consensus motif (23) frequently associated to an ACTACAN motif (39). The striking identity of this extended motif with SBS2 suggests a possible overlap between the RBPJ/ICN1 and ZNF143-binding events (Figure 5A). To test this assumption, we performed a ChIP-Seq experiment in the HPB-ALL cell line using an antibody directed against ICN1 (Supplementary Table S1). The data were compared with the ZNF143-ChIP-Seq data from the same cells. We identified that among the 1886 ZNF143- and 8467 ICN1-binding events located in promoters (Supplementary Table S1), 91.9% of the ZNF143-binding events are shared with those of the ICN1 protein (Figure 5B and Supplementary Figure S13). On average, the positions of the ICN1 and ZNF143 co-peaks coincided exactly (Figure 5C), suggesting the recognition of common sites by the two proteins. We also evidenced, as for ZNF143, that ICN1-binding events have a higher enrichment signal when associated to an SBS2 than an SBS1 site (t-test: \( P < 10^{-5} \)) (Figure 5D). By band shift assay, we showed that RBPJ binds specifically to SBS1 and SBS2 motifs, both containing the TTCCCA RBPJ-binding motif (Figure 5E).
Surprisingly, we observed that the RBPJ-binding was more efficient on the SBS2 motif, suggesting the possible involvement of the ACTACAN motif in recognition. Band shift assays performed on SBS1 or SBS2 motifs in the presence of ZNF143 and RBPJ revealed that the binding of the two proteins is mutually exclusive (Figure 5F). This observation is also supported by the results of an in vivo competition assay showing that over-expression of RBPJ provides a significant decrease of the ZNF143 occupation on target promoters common to ZNF143 and RBPJ (Figure 5G). Globally, this suggests that the ZNF143-ICN1 co-peaks observed by ChIP-Seq did not result from a simultaneous co-occupation of the DNA by both proteins. Finally, the analysis of ICN1-ChIP-Seq data also revealed that the MLT1J subfamily of retroelements alone was significantly enriched in ICN1-binding events (Supplementary Figure S12B).

**ZNF143, THAP11 and ICN1 levels directly modulate expression of common target genes**

The overlapping binding pattern of ZNF143, THAP11 and ICN1 led us to examine whether they play a direct role in the regulation of common target genes. To answer this question, we analysed the global or partial gene expression pattern after alteration of the level of the three factors. First, we performed global profiling of gene expression by RNA-Seq in cells depleted of or over-expressing ZNF143. ChIP assays with anti-β-catenin antibody were performed on induced FLP143-HA and FLPTHAP11-HA stable cell lines. Uninduced stable cell lines were used as control. THAP11-HA data are obtained from ChIP-qPCR experiment, amplifying a promoter region containing the binding site. Primer sequences are available on request. ZNF143-HA data are obtained from ChIP-Seq experiment, using read number values in regions amplified by qPCR for THAP11-HA, expanded to 200 bp. SBS1 sites and SBS2 sites correspond to promoters containing only the SBS1 motif or only the SBS2 motif, respectively. Fold enrichment in promoters was normalized to control regions located at 2 kb of tested promoters. The gene symbols listed are those of the genes closest to tested genomic region.
Figure 5. ICN1-binding regions co-localize with the ZNF143-binding events. (A) Sequence alignment of the RBPJ-binding site, SBS1 and SBS2 motifs. (B) Venn diagram depicting the overlap of ZNF143 and ICN1-binding events in HPB-ALL cells. (C) Overlap of ICN1 sites with ZNF143-binding sites in HPB-ALL cells. Average profile of ICN1 and ZNF143 ChIP-Seq signals around the centre of all overlapping peaks. (D) Box plots showing ChIP-Seq score (number of reads) distribution of ICN1-binding events associated to SBS1 or SBS2 motifs. (E) RBPJ in vitro binding assays on SBS1 and SBS2 motifs. Gel retardation assays were performed with $^{32}$P labelled double stranded fragment containing the indicated motif. The labelled probe was incubated in the presence (+) or absence (−) of RBPJ. Reactions in lanes 3, 7 and 4, 8 were performed in the presence of an excess of unlabelled specific (sp.) or unspecific (unsp.) competitors. The RBPJ–DNA complex is indicated by an arrow, the asterisk denotes an unrelated product. (F) The binding of RBPJ and ZNF143 is mutually exclusive in vitro. Gel retardation assays were performed with $^{32}$P labelled double-stranded fragment containing the SBS2 motif in the presence (+) or absence (−) of the indicated proteins. Arrows indicate the RBPJ–DNA and ZNF143–DNA complexes. (G) The binding of RBPJ and ZNF143 is mutually exclusive in vivo. ZNF143-ChIP-qPCR enrichment analysis on promoters bound by ZNF143 and Notch1. ZNF143-ChIP was performed on non-induced (ChIP reference) and induced FLP_RBPJ-HA cells expressing the RBPJ-HA (ChIP over-expression RBPJ) protein. ChIP enrichment was measured by qPCR using specific primers on promoter compared with negative regions located at 2 kb. Primers are available on request. A control ChIP was performed using non-specific IgG antibodies (ChIP control IgG). The gene symbols listed are those of the genes closest to the tested genomic region.
siRNA-mediated knockdown and over-expression efficiencies were confirmed at the protein (Figure 6A, lanes 1–6). After deep RNA sequencing, we examined the relative expression of the genes located at +/-2kb from a ZNF143 peak (Supplementary Table S2). The analysis revealed two major populations of differentially expressed target genes. The largest population (837 genes), as expected, was downregulated after ZNF143 knockdown and broadly upregulated after ZNF143 over-expression (Figure 6B, left part). The second, smaller population (386 genes), was unexpectedly upregulated after ZNF143 knockdown and mostly downregulated following its over-expression (Figure 6B, right part). The general tendency of the profiles suggests that on the full set of genes, ZNF143 acts mainly as an activator but can also be involved in the repression of transcription. An independent verification of the RNA-Seq data by RT-qPCR on a set of 51 genes activated or repressed after ZNF143 knockdown, recapitulated the majority of the findings (Supplementary Figure S14A). The genes positively regulated by ZNF143 (downregulated after ZNF143 knockdown) are primarily associated with ontology terms involved in transcription (mainly zinc-finger DNA-binding proteins), cell cycle regulation and chromatin organization (Supplementary Figure S14B, red bars). On the other hand, the negatively regulated genes (upregulated after ZNF143 knockdown) were only slightly enriched in genes involved in transcription processes (Supplementary Figure S14B, blue bars).

As we found THAP11 and ICN1 associated with ZNF143-binding motifs, we next analysed by RT-qPCR the effect of THAP11 and ICN1 knockdown on 51 ZNF143 target genes. After THAP11 knockdown in HeLa-S3 cells using specific siRNA (Figure 6D), 28 of the 51 genes were significantly upregulated and 4 of the 51 downregulated, in agreement with the major repressor function of the factor. After knockdown of ICN1 (Figure 6A, lanes 7–10), by γ-secretase inhibitor (GSI) treatment on HPB-ALL cells, 6 of the 51 genes were significantly upregulated and 20 of the 51 downregulated, in agreement with the major repressor function of the factor. After knockdown of ICN1 (Figure 6A, lanes 7–10), by γ-secretase inhibitor (GSI) treatment on HPB-ALL cells, 6 of the 51 genes were significantly upregulated and 20 of the 51 downregulated, in agreement with the major repressor function of the factor. After knockdown of ICN1 (Figure 6A, lanes 7–10), by γ-secretase inhibitor (GSI) treatment on HPB-ALL cells, 6 of the 51 genes were significantly upregulated and 20 of the 51 downregulated, in agreement with the major repressor function of the factor. After knockdown of ICN1 (Figure 6A, lanes 7–10), by γ-secretase inhibitor (GSI) treatment on HPB-ALL cells, 6 of the 51 genes were significantly upregulated and 20 of the 51 downregulated, in agreement with the major repressor function of the factor. After knockdown of ICN1 (Figure 6A, lanes 7–10), by γ-secretase inhibitor (GSI) treatment on HPB-ALL cells, 6 of the 51 genes were significantly upregulated and 20 of the 51 downregulated, in agreement with the major repressor function of the factor. After knockdown of ICN1 (Figure 6A, lanes 7–10), by γ-secretase inhibitor (GSI) treatment on HPB-ALL cells, 6 of the 51 genes were significantly upregulated and 20 of the 51 downregulated, in agreement with the major repressor function of the factor.

Promoters with associated SBS1 or SBS2 motifs carry different levels of RNA polymerase II and histone activation marks

The aforementioned data revealed that SBS1- and SBS2-mediated modulation of gene expression involves a competitive binding of different transcription factors on these motifs. We sought then to understand whether this differential occupation could be correlated with variations in Pol II occupancy and various histone H3 methylation and acetylation marks associated to activation or repression of transcription. It is well established that in the promoter of Pol II transcribed genes, the activating H3K4me3 and H3K27ac marks form two peaks on either side of the TSS, whereas the H3K36me3 mark accumulates within the transcribed region (40). Enhancers are commonly characterized by the presence of both H3K27ac and H3K4me1 histone marks (41). In contrast, the repressive marks H3K9me3 and H3K27me3 are associated to the inactive state of the chromatin (40). We first examined in HeLa-S3 cells the presence of H3K27ac and H3K4me1 around all ZNF143 peaks: ZNF143 peaks in promoters (+/-2 kb of a TSS) and ZNF143 peaks outside of promoters (Supplementary Figure S15). ZNF143 enriched regions outside of promoters do not co-localize with H3K27ac and H3K4me1 enhancer marks (Supplementary Figure S15C). On the contrary, peaks located in promoters are all highly enriched in ZNF143 and H3K27ac activation marks (Supplementary Figure S15B). Subsequently, we examined in HeLa-S3 cells the relation between histone H3 trimethylation and Pol II occupancy patterns on the ZNF143-binding events located near TSS. Heat maps representing the enrichment reveal a broad co-localization of ZNF143, Pol II and H3K4me3 modifications (Supplementary Figure S16A). Similarly, the H3K36me3 modification is found on the body of the genes associated to ZNF143-binding events (Supplementary Figure S16B). In contrast to H3K4me3, there was no detectable H3K9me3 and H3K27me3 modification associated to ZNF143-binding events (Supplementary Figure S16A). We observed a differential behaviour of loci associated to SBS1 or SBS2 motifs with respect to Pol II, H3K4me3 and H3K36me3 enrichments. Indeed, in opposition to ZNF143 enrichment (Figure 2D and Supplementary Figure S16C), the Pol II (Figure 6E), H3K4me3 (Figure 6F) and H3K36me3 enrichments (Supplementary Figure S16B) are higher on SBS1 than on SBS2-associated loci (t-test: P < 10^-5). Furthermore, the Pol II occupancy profile forms two peaks on each side of SBS2 loci, whereas a single one is observed for loci associated to SBS1 (Figure 6E). A similar behaviour of Pol II and H3K4me3 on SBS1 and SBS2 sites is observed for two different ChIP-Seq data sets from K562 and mES cells (Supplementary Figure S16D). The aforementioned observations suggest a distinct transcriptional control on SBS2 sites owing to the preferential binding of THAP11 on the SBS2 motif. We tested by ChIP-qPCR the Pol II occupation on several promoters containing SBS2 motifs after ZNF143 siRNA-mediated knockdown associated or not to THAP11 overproduction (Figure 6G). As expected, the results show a decrease of Pol II occupancy after ZNF143 knockdown, and the Pol II deficiency is amplified after THAP11 overexpression (Figure 6G).
Figure 6. ZNF143, THAP11 and ICN1 co-regulate the expression of target genes. (A) Western blots showing (i) knockdown of ZNF143 at 72 h post-transfection of HeLa cells with specific siRNA (lane 3) and siRNA control (lanes 1 and 2); (ii) over-expression of ZNF143 in FLP143 cells at 12 h (lane 5), 24 h (lane 6) post-induction, uninduced cells (lane 4); (iii) knockdown of ICN1 at 48 h and 72 h post-treatment of HPB-ALL cells with 500 nM of GSI (compound E) (lanes 9 and 10), DMSO treated (lane 8) and non-treated cells (lane 7). Tubulin was used as a loading control. (B) Bar chart depicting the RNA-Seq expression profiling of all genes located at +/−2 kb of a ZNF143 peak summit, after knockdown or over-expression of ZNF143. In grey and black are represented the fold change of expression 48 h after siRNA-mediated knockdown and 24 h after induced over-expression of ZNF143 compared with control, respectively. The genes are ordered decreasingly according their positive or negative fold change expression after ZNF143 knockdown. (C) Bar chart depicting the relative gene expression ratio determined by RT-qPCR for genes containing SBS1 or SBS2 sites, after ICN1 and ZNF143 knockdown compared with control. RNA was extracted 72 h post-transfection of HeLa-S3 cells with specific siRNAs targeting ZNF143 and from HPB-ALL cells 72 h post-treatment with 500 nM of GSI (compound E). Values represent the mean +/− SD of two or three independent experiments normalized to the GAPDH level. The dark grey line corresponds to a 0.5-fold expression ratio (D) Bar chart depicting the relative gene expression ratio determined by RT-qPCR for genes containing SBS1 or SBS2 sites, after THAP11 and ZNF143 knockdown compared with control. RNA was extracted 72 h post-transfection of HeLa-S3 cells with specific siRNAs targeting ZNF143 or THAP11. Values represent the mean +/− SD of two or three independent experiments normalized to the GAPDH level. The dark grey line corresponds to a 0.5-fold expression ratio (E) Average Pol II profile of Pol II and (F) Average H3K4me3 profile around the centre of ZNF143-binding events identified in HeLa cells at +/−2 kb from a TSS (Supplementary Table S4). All ZNF143-binding events, binding events containing only SBS1 or only SBS2, are indicated in black, dark grey and light grey, respectively. (G) RNA Pol II-ChIP-qPCR enrichment analysis on promoters bound by THAP11 and containing SBS2 sites. Pol II-ChIP was performed on cells treated with control siRNA (siCTRL), knockdown for ZNF143 (siZNF143) or knockdown for ZNF143 and over-expressing THAP11 (siZNF143 + THAP11-HA). ChIP enrichment was measured by qPCR using specific primers on promoter compared with negative regions located at 2 kb. Values represent the mean +/− SD of two or three independent experiments normalized to the GAPDH promoter levels. Primes are available on request. The gene symbols listed are those of the genes closest to the tested genomic region.
DISCUSSION

In this study, we obtained a set of >3000 genes that may be subject to a ZNF143-mediated regulation in both humans and mouse. These genes are globally highly enriched in terms related to transcription, cell cycle, RNA processing and translation processes, whereas cell signaling and cell development were under-represented. In addition, we observed as reported for PC3 cells (37) that the ZNF143 knockdown reduces significantly the cell proliferation rate in HeLa cells. These observations strongly suggest a particular role for this factor in cell growth regulation. We also report that both activation and repression of gene expression directed by ZNF143 occur in human cells.

Among the ZNF143 target genes, it is worth mentioning 114 KRAB zinc-finger transcription factors located in tandem on human chromosome 19. Considering the amino acid sequence similarity between the subdomain B of the KRAB box and the ncRNA transcription activation motif of ZNF143 (42), it is tempting to speculate that ZNF143 could originate from an ancient KRAB family member. A co-evolutionary link exists between retroelements and the presence of tandem zinc-finger genes (43). If we take into consideration the SBS2 site embedded in MLT11 retrotransposons, we can hypothesize that ZNF143 have diverged into a transcriptional regulator of more recent genes of the same family, generated by the successive invasion waves of genomes by KRAB zinc-fingers (44).

According to our results, ZNF76, the vertebrate paralog of ZNF143, occupies exactly the same binding sites as those recognized by ZNF143. We previously showed that ZNF76 is able to activate transcription (16), but other results have also identified ZNF76 as a transcriptional repressor (38). Having the same DNA-binding specificity as ZNF143, ZNF76 could have a different non-redundant function or it could play an auxiliary replacement function in particular situations.

As many ZNF143-binding events are located at proximity of ncRNA genes, this raises the question of the functionality of these events in ncRNA gene transcription. For the ZNF143 occupancy on 12 Pol III type 3 and 44 Pol II snRNA genes, it is very likely that the factor is involved in their transcriptional activation. Indeed, as very recently reported (45), all of these promoters contain the canonical sequence elements (35) and are bound by their proteins counterparts (46). Thirteen miRNA genes and a large number of snoRNA and scaRNA genes, found near ZNF143-binding events, are located in the first exon or in the first intron of coding or non-coding host genes. This opens the possibility that ZNF143 is involved in the expression of these ncRNAs by directing expression of the host gene. This is observed for MIR632, MIR659, MIR663, MIR933, MIR1292 and MIR3917 for which the host genes are deregulated in cells depleted or over-expressing ZNF143.

A major aspect of our study concerns the identification and the functional characterization of the SBS1 (18 bp) and SBS2 (25 bp) motifs associated to ZNF143-binding events. The additional ACTACAN sequence found in SBS2 has already been reported as part of one of the most represented motifs—the M4 motif—in gene promoters (47) and is not required for ZNF143 binding. It comes out from our study that the TBS ACTACAA TCC CAG is always physically linked to an overlapping ZNF143-binding site at the whole genome scale. We also evidenced, in vitro as well as in vivo, that the binding of ZNF143 and THAP11 on the SBS2 motif is mutually exclusive. These observations clearly indicate that the SBS2 subset of ZNF143-binding sites contains the set of TBSs. Moreover, we also demonstrated that the two proteins are both involved in the transcriptional regulation of the same set of genes. This gene expression modulation is achieved through a mutually exclusive binding on overlapping binding sites. As the result of the competitive binding of the two factors and to the main repressor activity of THAP11 (25), the promoters containing SBS2 sites have lower Pol II and H3K4/36me3 ChIP-Seq profiles. Nevertheless, unlike Pol II and H3K4me3 marks, ZNF143 shows a notably higher ChIP-Seq enrichment on SBS2 sites than on SBS1 sites. This observation supports the notion that SBS1- and SBS2-containing promoters are in different regulatory contexts.
The examination of SBS1 and SBS2 sites led us to consider another overlapping binding event with ICN1. Indeed, we demonstrated that in the T-lymphoblastic leukaemia cell line HPB-ALL, the constitutively over-expressed ICN1 protein is recruited in 27.6% of the ICN1-binding events via the TTCCCA sequence embedded in SBS1 and SBS2 motifs. In addition, we demonstrated that the RBPJ protein, which recruits ICN1, binds in vitro as well as in vivo to the SBS1 and SBS2 sites in an exclusive manner in the presence of ZNF143. Another recent study performed in a different T-lymphoblastic leukaemia cell line (CUTLL1) also highlighted that the ICN1-binding profile is partially overlapping ZNF143-binding sites (48). However, the authors of this study identified only an SBS2-type site as being bound by ZNF143. In contrast, our work performed on seven different human and mouse cell types clearly distinguishes the two motifs SBS1 and SBS2 as bound by ZNF143 and subjected to differential binding by two other unrelated transcription factors. As for ZNF143, our ChIP-Seq data on ICN1 showed higher tag enrichment for peaks associated to the SBS2 than to the SBS1 sites. This is consistent with our hypothesis that promoters containing either the SBS1 or SBS2 site are differently regulated. The gene expression analysis after GSI treatment in HPB-ALL cells confirmed the major activation function of ICN1 binding on both SBS1 and SBS2 sites.

We can infer from our results that ZNF143, THAP11 and ICN1/RBPJ are not only involved in a competitive binding on overlapping sites but are also involved in the transcriptional activity of adjacent genes. We revealed in this study that the genes targeted by ZNF143 are mainly involved in biological processes related to cell growth such as transcription, translation and cell cycle. Our observation is consistent with several reports concerning the role of THAP11 and ICN1. Indeed, the capacity of THAP11 to exert anti-differentiation effects in mES cells was originally suggested from THAP11 global dependent transcriptional repression of multiple genes that are either directly or indirectly involved in differentiation (27). In contrast, more recent results in mES cells showed that THAP11 can either activate or repress expression of its transcriptional target genes, essentially involved in the growth of embryonic stem cells (26). ICN1 is constitutively over-expressed in acute T-cell lymphoblastic leukaemia cells where it deregulates genes involved in cell growth processes (48,49). ZNF143, as shown in this study, is also involved in the regulation of genes involved in cell growth and has been shown to be important to maintain the undifferentiated state of mES cells (19). Taken together, it appears that ZNF143, THAP11 and ICN1 could together be important for the transcriptional modulation of genes involved in the proliferation of rapidly dividing cells, like cancer cells and embryonic stem cells. Therefore, this control is exerted via a competitive binding on overlapping binding sites.

All the observations and results taken together led us to propose the following model (Figure 7). The competitive and combinational binding of ZNF143, THAP11 and RBPJ/ICN1 on single or multiple SBS1 and SBS2 sites in promoters is directly involved in the transcriptional regulation of target genes. This is achieved by activation via ZNF143 and repression via THAP11 and RBPJ. Moreover, in HPB-ALL cells, the over-expressed ICN1 recruited by the repressor RBPJ competes for activation either with ZNF143 on SBS1 motifs or with THAP11 and ZNF143 on SBS2 motifs (Figure 7). Nevertheless, our model is restricted to certain genes being activated by ZNF143 and repressed by THAP11. Other genes, such as ALG14 and JUN, are negatively regulated by both THAP11 and ZNF143. This suggests an additional modulation mechanism that involves probably other partners acting positively or negatively on transcription. Such an additional partner could be ZNF76 that binds exactly the same motif as ZNF143. This extra layer of complexity should be further investigated.

Another interesting outcome of this work concerns the evolutionary history of SBS2 sites. As shown for several transcription factors (50), it emerged from our data that the very ancient family of MLT1J retro-transposons could be at the origin of the SBS2 sites. Furthermore, a correlation exists between KRAB zinc-finger gene expansions and retro-transposition waves (43). We thus speculate that ZNF143 originated from an ancient KRAB zinc-finger transcription factor family targeting at the time still active MLT1J retro-transposons. In the course of evolution, these events could have caused expansion of the repertoire of ZNF143, THAP11 and ICN1 regulated genes by spreading out the SBS2 sites in promoter regions.

SUPPLEMENTARY DATA

Supplementary Data are available at NAR Online: Supplementary Tables 1–7, Supplementary Figures 1–16 and Supplementary Materials and Methods.

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